



One Montvale Avenue
Stoneham, Massachusetts 02180
(781) 279-1675
FAX: (781) 279-1742

WARNING LETTER
NWE-DO-19-00W

February 1, 2000

VIA FEDERAL EXPRESS

Royal J. Bartrum, Jr., M.D.
Gifford Memorial Hospital
Post Office Box 2000
44 South Main Street
Randolph, VT 05060

RE: Inspection ID - 1139440005

Dear Dr. Bartrum,

We are writing to you because on January 26, 2000, your facility was inspected by a representative of the State of Vermont, acting in behalf of the Food and Drug Administration (FDA). This inspection revealed a serious regulatory problem involving the mammography at your facility.

Under a United States Federal law, the Mammography Quality Standards Act of 1992, your facility must meet specific requirements for mammography. These requirements help protect the health of women by assuring that a facility can perform quality mammography. The inspection revealed the following level 1 and level 2 findings at your facility:

Level 1: Phantom QC records were missing for 12 weeks for x-ray unit 2, [REDACTED] located in the mammography room.

Level 2: The radiologic technologist, [REDACTED], did not meet the continuing education requirement of having completed a minimum of 15 CEUs in mammography, in a 36 month period. No CEU's were completed.

The specific problems noted above appeared on your MQSA Facility Inspection Report, which was issued to your facility at the close of the inspection.

Because these conditions may be symptomatic of serious underlying problems that could compromise the quality of mammography at your facility, they represent a serious violation of the law which may result in FDA taking regulatory action without further notice to you. These actions include, but are not limited to, placing your facility under a Directed Plan of Correction, charging your facility for the cost of on-site monitoring, assessing civil money penalties up to \$10,000 for each failure to substantially comply with, or each day of failure to substantially comply with, the Standards, suspension or revocation of your facility's FDA certificate, or obtaining a court injunction against further mammography.

It is necessary for you to act on this matter immediately. Please explain to this office in writing within fifteen (15) working days from the date you received this letter:

- the specific steps you have taken to correct all of the violations noted in this letter;
- each step your facility is taking to prevent the recurrence of similar violations;
- equipment settings (including technique factors), raw test data, and calculated final results, where appropriate; and
- sample records that demonstrate proper record keeping procedures, if the findings relate to quality control or other records (Note: Patient names or identification should be deleted from any copies submitted).

Please submit your response to:

Michael J. Leal
U.S. Food & Drug Administration
120 Front Street, Suite 680
Worcester, MA 01608
(508) 793-0422
fax: (508) 793-0456

Finally, you should understand that there are many FDA requirements pertaining to mammography. This letter pertains only to findings of your inspection and does not necessarily address other obligations you have under the law. You may obtain general information about all of FDA's requirements for mammography facilities by contacting the Mammography Quality Assurance Program, Food and Drug Administration, P.O. Box 6057, Columbia, MD 21045-6057 (1-800-838-7715) or through the Internet at <http://www.fda.gov>.

If you have more specific questions about mammography facility requirements, or about the content of this letter, please feel free to contact Mr. Leal at (508) 793-0422.

Sincerely yours,



Michael R. Kravchuk
Acting District Director

cc:
Raymond N. McCandless
Director
Division of Occupational and Radiological Health
Department of Health
108 Cherry Street
P.O. Box 70
Burlington, VT 05402

Wilcox-Buchalla, R.N., M.B.A.
Director, Accreditation Programs
American College of Radiology
1891 Preston White Drive
Reston, Virginia 22091